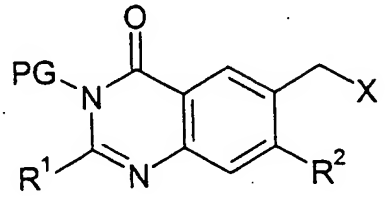
10/562112
JC10 Rec'd PCT/PTO 23 DEC 2005

AMENDMENTS TO THE CLAIMS:

This listing of claims will replace all prior versions, and listings, of claims in the application:

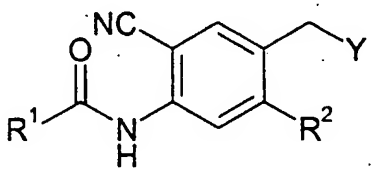
1 (original). A process for the preparation of a quinazolin-4-one derivative of formula (I):



(I)

where R^1 and R^2 are each independently hydrogen or methyl, PG is a protecting group and X is a leaving group;

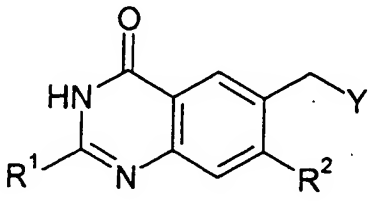
including the step of cyclization an amide of formula (II):



(II)

wherein R^1 and R^2 are as defined above and Y is a leaving group; or a protected derivative thereof;

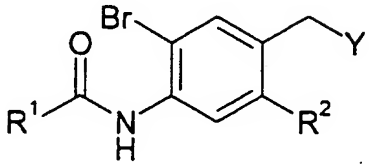
to form a quinazolin-4-one derivative of formula (III):



(III)

or a protected derivative thereof.

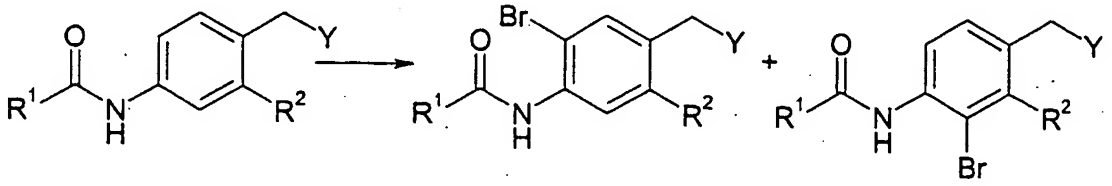
2 (original). A process as claimed in claim 1 wherein the amide of formula (II) is made by reacting a compound of formula (IV):



(IV)

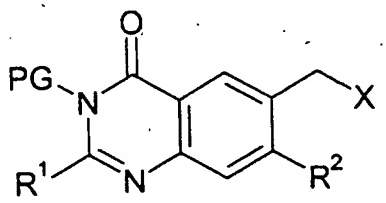
with a cyanide reagent.

3 (original). A process as claimed in claim 2 wherein the compound of formula (IV) is made by a regioselective bromination step from a compound of formula (V) using the reaction step:



(V) (IV) (IVA)

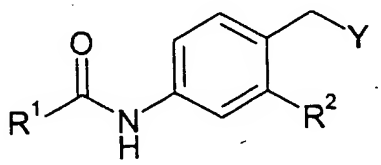
4 (original). A process for the preparation of a quinazolin-4-one derivative of formula (I):



(I)

where R^1 and R^2 are each independently hydrogen or methyl, PG is a protecting group and X is a leaving group;

including the step of brominating a compound of formula (V):

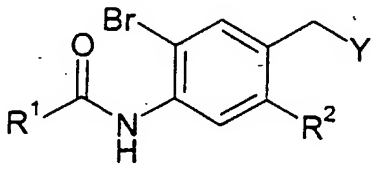


(V)

wherein R^1 and R^2 are as defined above and Y is a leaving group;

or a protected derivative thereof;

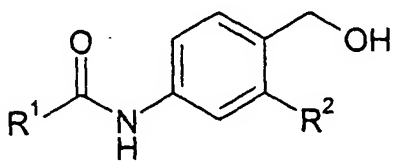
to form a compound of formula (IV)



(IV)

or a protected derivative thereof.

5 (original). A process as claimed in claim 4 wherein the compound of formula (V) is made by derivatization of an alcohol of formula (VI):

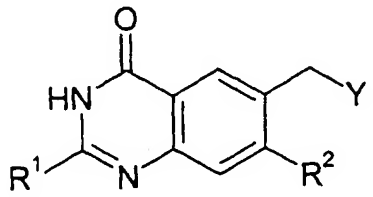


(VI)

6 (currently amended). A process as claimed in ~~any preceding claim~~ claim 1 wherein at least one of R^1 and R^2 is methyl.

7 (currently amended). A process as claimed in ~~any~~ claim 6 wherein R^1 and R^2 are both methyl.

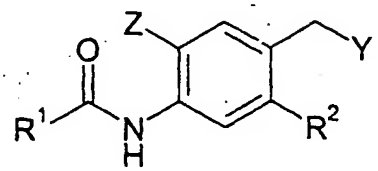
8 (original). A quinazolin-4-one derivative of formula (III):



(III)

where R^1 and R^2 are each independently hydrogen or methyl, and Y is a C_{1-4} acyloxy group or benzoyloxy.

9 (original). An amide of formula (VIII):



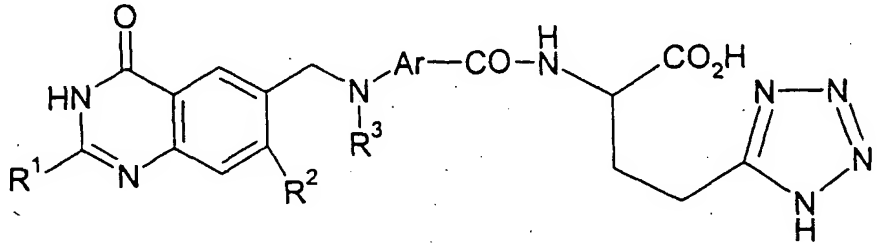
(VIII)

wherein R^1 and R^2 are each independently hydrogen or methyl, Y is a C_{1-4} acyloxy group or benzoyloxy and Z is Br or CN.

10 (currently amended). A compound as claimed in claim 8 ~~or claim 9~~ wherein at least one of R^1 and R^2 is methyl.

11 (original). A compound as claimed in claim 10 wherein R^1 and R^2 are both methyl.

12 (currently amended). A process as claimed in ~~any one of claims 1 to 7~~ claim 1 wherein the process is used to prepare a quinazoline-4-one of formula (IX):



(IX)

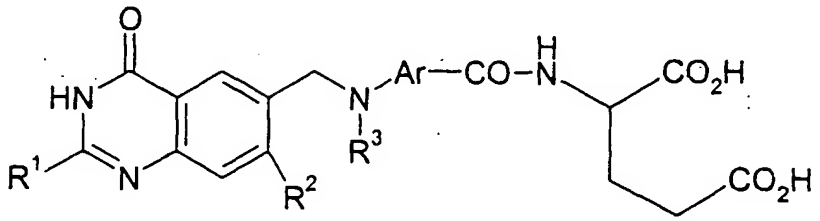
wherein R^1 and R^2 are each independently hydrogen or methyl;

R^3 hydrogen, C_{1-4} alkyl, C_{3-4} alkenyl, C_{3-4} alkynyl, C_{2-4} hydroxyalkyl C_{2-4} halogenoalkyl or C_{1-4} cyanoalkyl;

and Ar is phenylene, thiophenediyl, thiazolediyl, pyridinediyl or pyrimidinediyl which may optionally bear one or two substituents selected from halogeno, hydroxy, amino, nitro, cyano, trifluoromethyl, C_{1-4} alkyl and C_{1-4} alkoxy;

or a pharmaceutically-acceptable salt or ester thereof.

13 (currently amended). A process as claimed in ~~any one of claims 1 to 7~~ claim 1 wherein the process is used to prepare a quinazoline-4-one of formula (X):



(X)

wherein R^1 and R^2 are each independently hydrogen or methyl;

R^3 hydrogen, C_{1-4} alkyl, C_{3-4} alkenyl, C_{3-4} alkynyl, C_{2-4} hydroxyalkyl C_{2-4} halogenoalkyl or C_{1-4} cyanoalkyl;

and Ar is phenylene, thiophenediyl, thiazolediyl, pyridinediyl or pyrimidinediyl which may optionally bear one or two substituents selected from halogeno, hydroxy, amino, nitro, cyano, trifluoromethyl, C_{1-4} alkyl and C_{1-4} alkoxy;

or a pharmaceutically-acceptable salt or ester thereof.